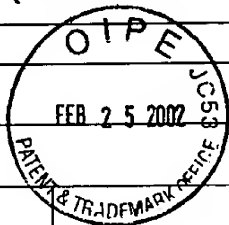


<b>FORM PTO-1449</b>		<b>ATTY. DOCKET NO.</b> 30727.001-IP1	<b>SERIAL NO.</b> 09/518,501
<b>LIST OF PATENTS AND OTHER ITEMS FOR APPLICANT'S INFORMATION DISCLOSURE STATEMENT</b>  <i>HH</i> (Use several sheets if necessary)		<b>APPLICANT:</b> METABASIS THERAPEUTICS, INC.	
		<b>FILING DATE:</b> March 5, 1999	<b>GROUP:</b> 1800



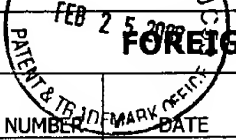
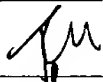

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U.S. PATENT DOCUMENTS							
EXAMINER INITIAL		DOCUMENT NUMBER	DATE	NAME	CLASS	CLASS	FILING DATE
<i>TUM</i>	AA	3,018,302	01/23/62	Bielefeld, et al.	260	461	—
<i>TUM</i>	AB	5,157,027	10/20/92	Biller, et al.	514	167	—
<i>TUM</i>	AC	5,658,889	8/19/97	Gruber, et al.	514	43	—

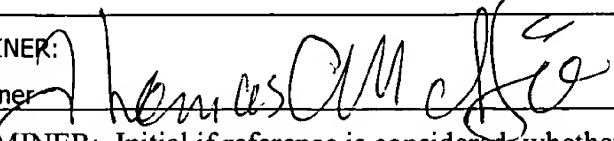
FOREIGN PATENT DOCUMENTS								
EXAMINER INITIAL		DOCUMENT NUMBER	DATE	COUNTRY	CLASS	SUB CLASS	TRANSLATION	
							YES	NO
<i>TUM</i>	AD	0 002 062 A1	30.05.79	EP (and English language Abstract)				
<i>TUM</i>	AE	0 072 531 A1	23.02.83	EP (and English language Abstract)				
<i>TUM</i>	AF	0 158 057 A	16.10.85	EP (U.K. English language translation)				
	AG	0 180 276 A1	28.12.88	EP				
	AH	0 261 283 A1	22.09.86	EP				
	AI	0 161 955 A1	21.11.85	EPO				
	AJ	0 338 372 A2	25.10.89	EPO				
	AK	0 353 692 A2	07.02.90	EPO				
	AL	0 481 214 B1	22.04.92	EPO				
	AM	492 738	30.06.70	CH (and English language Abstract)				
<i>✓</i>	AN	987,378	17.09.70	UK				

<b>EXAMINER:</b> <i>James M. Chis</i>	<b>DATE CONSIDERED:</b> <i>7/8/02</i>
EXAMINER: Initial if reference is considered, whether or not citation is in conformance with MPEP 609; Draw line through citation if not in conformance and not considered. Include a copy of this form with next communication to applicant	

<b>FCRM PTO-1449</b>  <b>LIST OF PATENTS AND OTHER ITEMS FOR APPLICANT'S INFORMATION DISCLOSURE STATEMENT</b>  (Use several sheets if necessary)	<b>ATTY. DOCKET NO.</b> 30727.001 [REDACTED] IP1	<b>SERIAL NO.</b> 09/518,501
	<b>APPLICANT:</b> METABASIS THERAPEUTICS, INC.	
	<b>FILING DATE:</b> March 5, 1999	<b>GROUP:</b> 1614

<div style="text-align: center;">  <b>FOREIGN PATENT DOCUMENTS</b> </div>								
EXAMINER INITIAL		DOCUMENT NUMBER	DATE	COUNTRY	CLASS	SUB CLASS	TRANSLATION YES	NO
 	AO	35 12 781 A1	10.04.85	DE (and English language U.S. Patent No. 4,952,740)				
	AP	96/01267	18.01.96	WO				
	AQ	97/03679	06.02.97	WO				
	AR	98/09668	12.03.98	WO				
	AS	98/39342	11.09.98	WO				
	AT	98/39343	11.09.98	WO				
	AU	98/39344	11.09.98	WO				

OTHER DOCUMENTS (Including Author, Title, Date, Pertinent Pages, etc.)		
 	AV	"Patent Abstracts of Japan," Vol. 1998, No. 1, 30 January 1998 & JP 09241284 A (Yamishata Koji, et al.; Nippon Soda Co. Ltd), 16 September 1997
	AW	DeLombaert, et al., "N-Phosphonomethyl Dipeptides and Their Phosphonate Prodrugs, a New Generation of Neutral Endoprtidase (NEP, EC 3.4.24.11) Inhibitors, <u>J. Med. Chem.</u> 37: 498-511 (1994)
	AX	Edmunson, et al., "Cyclic Organophosphorus Compounds. Part 23. Configurational Assignments in the 4-Phenyl-1,3,2λ5-dioxaphosphorinane Series. X-Ray Molecular Structure of <i>cis</i> -2-Benzylamino-4-phenyl-1,3,2-dioxaphosphorinane 2-Oxide," <u>J. Chem. Res. Synop.</u> , 5: 122-123 (1989)
	AY	Farquhar, et al., "5'-4-(Pivaloyloxy)-1, 3, 2-dioxaphosphorin an -2-yl]-2'-deoxy-5-fluorouridine: a membrane-permeating prodrug of 5-fluoro-2'-deoxyuridylic acid (FDUMP)" <u>J. Med. Chem.</u> 38:488-495 (1995)
	AZ	Farquhar, et al., "Biologically Reversible Phosphate-Protective Groups," <u>J. Pharm. Sci.</u> 72(3): 324-325 (1983)

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SERIAL NO.

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**OTHER DOCUMENTS (Including Author, Title, Date, Pertinent Pages, etc.)**

BA	Farquhar, et al., "Synthesis and Biological Evaluation of 9-[5'-(2-Oxo-1,3,2-oxazaphosphorinan-2-yl)-β-D-arabinosyl]adenine and 9-[5'-(2-Oxo-1,3,2-dioxazaphosphorinan-2-yl)-β-D-arabinosyl]adenine: Potential Neutral Precursors of 9-[β-D-Arabinofuranosyl]adenine 5'-Monophosphate," <u>J. Med. Chem.</u> 28: 1358-1361 (1985)
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BD	Hillers, et al., "Analogues of pyrimidinemono- and polynucleotides. IV. Phosphates of 1-(1,4-dihydroxy-2-pentyl)thymine and 1-(1,3-dihydroxy-2-propyl) uracil." <u>Khim Geterotski Soedin</u> 5:678-683 (1978). Chem Abst. v 89 no 17; abst no 146864u
BE	Hunston, et al., "Synthesis and Biological Properties of Some Cyclic Phosphotriesters Derived from 2'-Deoxy-5-fluorouridine," <u>J. Med. Chem.</u> 27: 440-444 (1984)
BF	Kryuchkov, et al., <u>Izv. Akad. Nauk SSSR, Ser. Khim.</u> 6: 1201-1248 (1987)
BG	Lok, et al., "Neurotoxicity associated with adenine arabinoside monophosphate in the treatment of chronic hepatitis B virus infection," <u>J. Antimicrob. Chemotherap.</u> 14: 93-99 (1984)
BH	Ludeman, et al., "Synthesis and Antitumor Activity of Cyclophosphamide Analogues. 4. Preparation, Kinetic Studies, and Anticancer Screening of "Phenyl ketophosphamide" and Similar Compounds Related to the Cyclophosphamide Metabolite Aldophosphamide," <u>J. Med Chem.</u> 29, 716-727 (1986)
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BJ	Meyer, et al., "2"-O'-Acyl-6-thioinosine Cyclic 3', 5'-Phosphates as Prodrugs of Thioinosinic Acid," <u>J. Med. Chem.</u> 22: 811-815 (1979)
BK	Neidlein, et al., "Mild Preparation of 1-Benzyloxyiminoalkylphosphonic Dichlorides: Application to the Synthesis of Cyclic Phosphonic Diesters and Cyclic Monoester Amides," <u>Heterocycles</u> 35: 1185-1203 (1993)

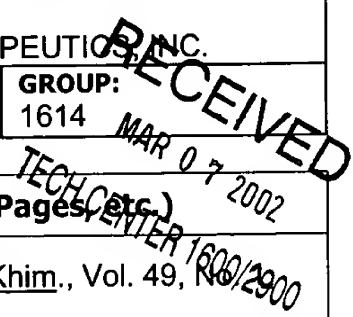
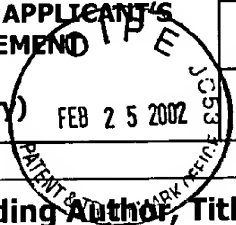
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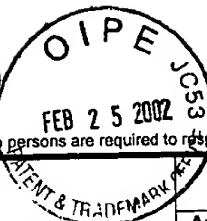
<b>FORM PTO-1449</b>		<b>ATTY. DOCKET NO.</b> 30727.001 IP1	<b>SERIAL NO.</b> 09/518,501
<b>LIST OF PATENTS AND OTHER ITEMS FOR APPLICANT'S INFORMATION DISCLOSURE STATEMENT</b>  (Use several sheets if necessary)		<b>APPLICANT:</b> METABASIS THERAPEUTICS, INC.	
		<b>FILING DATE:</b> March 5, 1999	<b>GROUP:</b> 1614



OTHER DOCUMENTS (Including Author, Title, Date, Pertinent Pages, etc.)		
BL	Nifant'ev, et al., "1,3,2, - Diazaphosphorinanes", <u>Zh. Obshch. Khim.</u> , Vol. 49, 1979, Pages 53-61.	
BM	Nifant'ev, et al., "Synthesis and Structure of Some Stable Phospholane-Phospholanes," <u>Phos. Sulfur &amp; Silicon</u> 113, 1-13 (1996)	
BN	Predvoditelev D. , et al., "Glycero-2-hydroxymethylene phosphates" <u>Journal of Organic Chemistry of the USSR (English Translation</u> 13:1489-1492 (1977))	
BO	Predvoditelev, D. et al., "Synthesis of lipids and their models on the basis of glycerol alkylene phosphites. V. Cyclic phosphatidylglycerol and phosphatidylhydroxyhomocholine" <u>Journal of Organic Chemistry of the USSR (English Translation</u> 17:1156-1165 (1981)	
BP	Shaw & Cundy, "Biological Screens of PMEA Prodrugs," <u>Pharm. Res.</u> 10 (Suppl) S-294 (1993)	
BQ	Shih, et al., "Preparation and Structures of 2-Dimethylamino-4-phenyl-1,3,2-dioxaphosphorinane-2-oxides," <u>Bull. Inst. Chem. Acad. Sin.</u> 41: 9-16 (1994)	
BR	Starrett, et al., "Synthesis, Oral Bioavailability Determination, and <i>in Vitro</i> Evaluation of Prodrugs of the Antiviral Agent 9-[2-(Phosphonomethoxy)-ethyl]adenine (PMEA)," <u>J. Med. Chem.</u> 37: 1857-1864 (1994)	
BS	Yamanaka, et al., "Metabolic Studies on BMS-200475, a New Antiviral Compound Active against Hepatitis B Virus," <u>Antimicrob. Agents Chemoth.</u> 43, 190-193 (1999)	
BT	Zon, et al., "4 Cyclophosphamide Analogues," <u>Prog. Med. Chem.</u> 19: 205-246 (1982)	

<b>EXAMINER:</b> Examiner	<b>DATE CONSIDERED:</b>
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# INFORMATION DISCLOSURE STATEMENT BY APPLICANT

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Sheet 1 of 2

Application Number 09/518,501  
Filing Date March 3, 2000  
First Named Inventor Erion  
Group Art Unit 1624  
Examiner Name T. McKenzie  
Attorney Docket Number 030727.0013.CIP1

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## U.S. PATENT DOCUMENTS

Examiner Initials*	Cite No. <sup>1</sup>	U.S. Patent Document		Name of Patentee or Applicant of Cited Document	Date of Publication of Cited Document MM-DD-YYYY	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear
		Number	Kind Code <sup>2</sup> (if known)			

## FOREIGN PATENT DOCUMENTS

Examiner Initials	Cite No. <sup>1</sup>	Foreign Patent Document			Name of Patentee or Applicant of Cited Document	Date of Publication of Cited Document MM-DD-YYYY	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear	T <sup>6</sup>
		Office <sup>3</sup>	Number <sup>4</sup>	Kind Code <sup>5</sup> (if known)				

## NON PATENT LITERATURE DOCUMENTS

Examiner Initials	Cite No. <sup>1</sup>	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	T <sup>2</sup>
	AA	Coppi, et al., "Lewis Acid Mediated Condensation of Alkenols and Aldehydes. A Selective Synthesis of Tetrahydropyrans and Oxepanes," J. Org. Chem., Vol. 53, No. 4, 911-913 (1988)	
	AB	Lohr et al., "Targeted chemotherapy by intratumor injection of encapsulated cells engineered to produce CYP2B1, and ifosfamide activating cytochrome P450," Gene Therapy, 5, 1070-1078 (1988).	
	AC	Bijsterbosch, et al., "Disposition of the Acyclic Nucleoside Phosphonate (S)-9-(3-Hydroxy-2-Phosphonylmethoxypropyl)Adenine," Antimicrobial Agents and Chemotherapy, Vol. 42, p. 1146-1150 (May 1998).	
	AD	de Waziers, et al., "Cytochrome P 450 Isoenzymes, Epoxide Hydrolase and Glutathione Transferases in Rat and Human Hepatic and Extrahepatic Tissues1," The Journal of Pharmacology and Experimental Therapeutics, Vol. 253, No.1, p. 387-394 (1989).	

Examiner Signature

[Signature]

Date

Considered

2/12/03

\*EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609. Draw line through citation is not in conformance and not considered. Include copy of this form with next communication to applicant.

<sup>1</sup> Unique citation designation number. <sup>2</sup> See attached Kinds of U.S. Patent Documents. <sup>3</sup> Enter Office that issued the document, by the two-letter code (WIPO Standard ST.3). <sup>4</sup> For Japanese patent documents, the indication of the year of the reign of the Emperor must precede the serial number of the patent document. <sup>5</sup> Kind of document by the appropriate symbols as indicated on the document under WIPO Standard ST. 16 if possible. <sup>6</sup> Applicant is to place a check mark here if English language Translation is attached.

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**Complete if Known**

<b>Application Number</b>	09/518,501
<b>Filing Date</b>	March 5, 1999
<b>First Named Inventor</b>	Erion
<b>Group Art Unit</b>	1624
<b>Examiner Name</b>	T. McKenzie
<b>Attorney Docket Number</b>	030727.0013.CIP1

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Sheet	2	of	2
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<sup>1</sup> Unique citation designation number. <sup>2</sup> See attached Kinds of U.S. Patent Documents. <sup>3</sup> Enter Office that issued the document, by the two-letter code (WIPO Standard ST.3). <sup>4</sup> For Japanese patent documents, the indication of the year of the reign of the Emperor must precede the serial number of the patent document. <sup>5</sup> Kind of document by the appropriate symbols as indicated on the document under WIPO Standard ST. 16 if possible. <sup>6</sup> Applicant is to place a check mark here if English language Translation is attached.

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30727.0013.CIP1SERIAL NO.  
09/518,501LIST OF PATENTS AND OTHER ITEMS FOR APPLICANT'S  
INFORMATION DISCLOSURE STATEMENTAPPLICANT:  
METABASIS THERAPEUTICS, INC.FILING DATE:  
March 5, 1999GROUP:  
1614

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## U.S. PATENT DOCUMENTS

EXAMINER INITIAL		DOCUMENT NUMBER	DATE	NAME	CLASS	SUB CLASS	FILING DATE
TC	AA	3,018,302	01.23.62	Bielefeld, et al.			
	AB	5,157,027	10/20 92	Biller, et al.			
	AC	5,658,889	8/19/97	Gruber, et al.			

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	AR	98/09668 A	12.03.98	WO			
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	AT	98/39343	11.09.98	WO			
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AV	"Patent Abstracts of Japan," Vol. 1998, No. 1, 30 January 1998 & JP 09 241284 A (Yamishata Koji; Nippon Soda Co. Ltd), 16 September 1997
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## OTHER DOCUMENTS (Including Author, Title, Date, Pertinent Pages, etc.)

BL	Nifant'ev, et al., "1,3,2, - Diazaphosphorinanes", <u>Zh. Obshch. Khim.</u> , Vol. 49, No. 1, 1979, Pages 64-74 (and English version as translated in corresponding English language publication)
BM	Nifant'ev, et al., "Synthesis and Structure of Some Stable Phospholane-Phospholanes," <u>Phos. Sulfur &amp; Silicon</u> 113, 1-13 (1996)
BN	Predvoditelev D. , et al., "Glycero-2-hydroxymethylene phosphates" <u>Journal of Organic Chemistry of the USSR (English Translation</u> 13:1489-1492 (1977))
BO	Predvoditelev, D. et al., "Synthesis of lipids and their models on the basis of glycerol alkylene phosphites. V. Cyclic phosphatidylglycerol and phosphatidylhydroxyhomocholine" <u>Journal of Organic Chemistry of the USSR (English Translation</u> 17:1156-1165 (1981)
BP	Shaw & Cundy, "Biological Screens of PMEA Prodrugs," <u>Pharm. Res.</u> 10 (Suppl) S24 (1993)
BQ	Shih, et al., "Preparation and Structures of 2-Dimethylamino-4-phenyl-1,3,2-dioxaphosphorinane-2-oxides," <u>Bull. Inst. Chem. Acad. Sin.</u> , 41: 9-16 (1994)
BR	Starrett, et al., "Synthesis, Oral Bioavailability Determination, and <i>in Vitro</i> Evaluation of Prodrugs of the Antiviral Agent 9-[2-(Phosphonomethoxy)-ethyl]adenine (PMEA)," <u>J. Med. Chem.</u> 37: 1957-1864 (1994)
BS	Yamanaka, et al., "Metabolic Studies on BMS-200475, a New Antiviral Compound Active against Hepatitis B Virus," <u>Antimicrob. Agents Chemoth.</u> 43, 190-193 (1999)
BT	Zon, et al., "4 Cyclophosphamide Analogues," <u>Prog. Med. Chem.</u> 19: 205-246 (1982)

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